1	IN THE UNITED STATES DISTRICT COURT FOR THE SOUTHERN DISTRICT OF WEST VIRGINIA AT CHARLESTON
3	TRANSCRIPT OF PROCEEDINGS
4	
5	x
6	: THE CITY OF HUNTINGTON, : CIVIL ACTION : NO. 3:17-cv-01362
7	Plaintiff, :
8	vs. :
9	AMERISOURCEBERGEN DRUG : CORPORATION, et al., :
10	: Defendants. :
12	: x
13	CABELL COUNTY COMMISSION, : CIVIL ACTION : NO. 3:17-cv-01665
14	Plaintiff, :
15	vs. :
16	AMERISOURCEBERGEN DRUG : CORPORATION, et al., :
17	: Defendants. :
18	: x
19	
20	VIDEO PRE-TRIAL CONFERENCE
21	BEFORE THE HONORABLE DAVID A. FABER
22	SENIOR UNITED STATES DISTRICT JUDGE
23	
24	April 14, 2021
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    Proceedings recorded by mechanical stenography; transcript
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    produced by computer.
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## PROCEEDINGS

THE COURT: All right. This is another pre-trial conference and motions hearing in the City of Huntington against AmerisourceBergen and others case.

I assume all the attorneys have given your appearances to the court reporter. And she's nodding "yes" and I hear no objection to that.

So the first thing I have on the list, I thought we'd -- I should say something about the logistics. I know there have been a lot of discussions and things about that.

Can you all hear me?

(All counsel indicated an affirmative response.)

THE COURT: All right. The Centers for Disease

Control limit suggests a cap of 30 people in the courtroom.

If we have five attorneys per party, which I believe there

have been discussions to allow, that's 25 people. The court

family will have at least five members. That's 30. Then,

of course, we'll have a witness and, in some cases, a

witness counsel. So that fills up the courtroom where the

trial is actually taking place.

As I've indicated before, I'm not going to allow a public feed on the -- I'm not going to allow any public feed. I will allow public feed for the final pre-trial.

I'm concerned about the obligation to grant the public and the press access to the trial. And since there's not going

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to be a public feed and there's so many persons involved,
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    that's difficult. But I think we can handle it.
          We're going to provide an overflow courtroom which will
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 4
     seat about 25 people I believe. And we can arrange a
5
     separate overflow courtroom with a capacity of about 35 more
     if, if we need it. Hopefully that will do it.
 6
7
          I read this morning the memo from Mr. Ruby suggesting
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    procedures, and I'm in agreement with everything in that.
9
    And hopefully that will give everybody guidance on, on the
10
     logistics.
11
          The first motion I have for hearing is the plaintiffs'
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    motion to compel. This is ECF Number 1211 from the court
13
     file. I understand Mr. Fuller for the plaintiffs and Ms.
14
    Wicht for the defendants will argue this.
15
          I'm informed that Ms. Wicht's monitor is not -- she's
16
    not getting any video now. The Court won't be able to see
17
    her.
18
          But I think we can hear you, Ms. Wicht, and we'll make
     the best of that situation.
19
20
          Mr. Fuller, do you want to go ahead?
21
               MR. FULLER: Yes, Your Honor, if it may please the
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    Court.
23
               THE COURT: Yes.
24
               MR. FULLER: Judge, this motion focuses around the
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sales and data of Cardinal Health not just to CT2 or West

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Virginia but across the country. And the, the information has been provided via the ARCOS database which Your Honor has heard some argument about that and Mr. McCann's working with that. Mr. McCann will provide some opinions and, and additional information related to that ARCOS database.

But all that information is is the defendants' sales data. And what I've tried to do with the 1006 that I provided to the defendants is simplify it because you're dealing with millions upon millions of lines of data or pills.

The 1006 basically sums up the Oxycontin and the hydrocodone -- excuse me -- oxycodone and hydrocodone that was delivered into each state. And there's significant reasons why that's important in this litigation because one of the things the defendants are looking for is what's unusual; unusual size, pattern, and frequency.

And what this data demonstrates is it demonstrates that while West Virginia is getting a significant number of pills, it outweighs what is going into other states and which would demonstrate what is unusual.

So, for example, just one for the Court, if we take a look at Texas, Texas is about 15 times larger in population. Yet, West Virginia got more oxycodone pills over the same time frame than Texas did which would certainly speak to something that is going on that is unusual.

Now, what the motion asks for, Judge, is not for you to admit the 1006. The defendants make some reference to that's what we're seeking. We're not. They've -- and, quite frankly, I've worked very well with Ms. Wicht and the other Cardinal counsel and we've been able to work a lot of issues out. But the 1006, they tell me they object to it. They won't specify why they object to it.

And then based on the stipulation that's part of the record in this case with Cardinal, I said, all right, well, give me a 30(b) witness that can tell me how the volume of pills going to each state from the time frame of '06 to '14, or just tell me how many pills you believe you've shipped into each state.

THE COURT: Let me interrupt you.

MR. FULLER: Yes, Your Honor.

THE COURT: The problem I have with this is what authority do I have to order the parties to do something that's within their discretionary management of their case?

Do you have any authority that says I can do this?

MR. FULLER: Well, Judge, I'm relying on the stipulation. The defendants are going to say, well, this isn't their data. Well, it is their sales data. And the stipulation requires if they're objecting to the admissibility of something that we have the opportunity to cure that before trial. And that's all I'm trying to do,

Judge.

THE COURT: I understand all that. But do you have any authority for what you want me to do? You're asking me to compel the defendants to stipulate here and I don't -- I question whether I have any authority to do that.

That's sticking my nose into, into their trial tactics and trial preparation. And there may well be a reason why they're taking this action. And, and I need some authority to persuade me that I can do this if I'm going to do it.

MR. FULLER: Sure, Judge, if that was what we were asking for. And that's not what we're seeking, Your Honor.

What I'm simply seeking is to require them to provide me a 30(b) who can give me the evidence that I need in a form that's admissible, or they can just disclose to us what they believe their sales data is.

The motion does not seek to have you require them to stipulate. It does seek to have you order them to allow me to correct any deficiencies that they are objecting to.

So I -- and I want to make that abundantly clear.

We're not asking you to force them to stipulate because I believe Your Honor is correct. You don't have the authority to do that. That's their prerogative.

But the Court does have the authority to require them to give me a 30(b) so I can correct the deficiencies that they perceive with the 1006.

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               THE COURT: Well, you ask in your motion for an
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     order compelling Cardinal Health, quote, to provide the
 3
     complete basis for refusing to stipulate to the accuracy of
    plaintiffs' request. So --
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 5
              MR. FULLER: Yes, Judge. I think -- I'm sorry.
 6
               THE COURT: So if, if I make them provide the
    basis for refusing to stipulate, that's, in effect, ordering
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8
     them to stipulate, isn't it? I mean, it's just one step
9
     away from that.
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               MR. FULLER: Well, Judge, I think it's requiring
11
     them to tell us what is the -- what they believe is
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     deficient with it. And I think also in our "wherefore"
13
     clause we ask that in the alternative they be ordered to
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    provide us what they believe their sales numbers are into
15
     each state or to give us a 30(b) that can testify based on
16
     the company's knowledge of what their sales numbers are into
17
     each state.
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               THE COURT: Well, why can't you just get a 30(b)
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    without the Court's intervention? Is that, is that
20
     inconsistent with the scheduling orders and so forth?
21
               MR. FULLER:
                            That, that is, Your Honor. That is.
22
    And the reason I'm seeking it -- and we attached the
23
     stipulation that we entered into with Cardinal on certain
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     issues -- is that that stipulation, when they object to
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     something being admissible, they stipulated that they would
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provide us even a deponent if that's what we needed to cure
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     the defects. And that's simply what we're trying to do.
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 3
         We're not asking this Court to force them to stipulate
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    because I think that would be absolutely improper. But the
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    Court could require them to provide a 30(b); or I think the
 6
    easiest way, Judge, is just say, hey, tell them what you
7
    believe the distribution numbers are into each state. And
8
     then I think we could work it out.
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               THE COURT: So, so basically -- I mean, I'm a
10
     little bit -- basically, what you're asking for is for me
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     to, in a very limited method, reopen discovery so you can
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     take this 30(b). Is that right?
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               MR. FULLER: Or if they want to just produce to us
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    what they believe the sales numbers are into each state, I
15
     think that would actually be the easiest, Judge. But, yes,
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     Your Honor, that's, that's what we're requesting.
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               THE COURT: And if, if you take the 30(b), you
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    would be after the information that you've just said you
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     thought they should voluntarily provide. Is that right?
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              MR. FULLER: Yes, Your Honor.
21
               THE COURT: Okay. Let me hear from Ms. Wicht.
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THE COURT: Okay. Let me hear from Ms. Wicht. I hope you can hear all this and I'll hear from you now. And if there's any problem with anybody hearing you, let me know. Apparently we've got a partial breakdown in technology here.

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MS. WICHT: Good morning, Your Honor. Thank you
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    very much. I am able to see and I'm actually able to see
     and hear everybody. I hope that everyone can hear me.
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               THE COURT: Well, I can certainly hear you and I
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    don't see anybody objecting on the screen here. So I
 6
    assume --
7
              MS. WICHT: Okay.
8
               THE COURT: And you're coming through loud and
9
     clear in the courtroom.
10
              MR. FULLER: We can hear fine, Judge.
11
              MS. WICHT: Thank you, Your Honor.
12
         Thank you, Mr. Fuller. I apologize for the technical
13
    difficulties.
14
               THE COURT: Can you hear her, Mr. Fuller? You're
15
    the key man here.
16
              MS. WICHT: I can.
17
              MR. FULLER: Yes, Your Honor. I can hear her
18
     clearly.
19
               THE COURT: All right. Go ahead, please.
20
               MS. WICHT: Thank you, Your Honor.
21
         What plaintiffs are actually seeking in this motion,
22
    Your Honor, is an order of the Court to force Cardinal
23
    Health to stipulate to the accuracy of the work of
24
    plaintiffs' expert.
25
         There's nothing in the Federal Rules or the prior
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stipulations of the parties that requires Cardinal Health to agree with plaintiffs' expert, nor to provide testimony after the expert's work was presented in an untimely fashion, I might add, to identify their agreements or disagreements with the expert.

Mr. Fuller and the plaintiffs in their papers and in their argument today framed this as being an issue about Cardinal Health's data, but that's not correct. That's not what's been presented in these charts.

What's presented in the charts is that Cardinal Health submitted distribution data to DEA through the ARCOS system.

DEA, presumably in the normal course of its operations, made changes to that data, including adding 22 fields of information that were generated and calculated by DEA, including the principal one that's featured on plaintiffs' summary which is dosage units.

DEA then produced that data to plaintiffs in this litigation. And plaintiffs' expert, Dr. McCann, conducted an analysis that resulted in other changes to the data that he deemed to be necessary to make it accurate in his opinion.

So that's the information that plaintiffs have summarized in the charts that they've presented. So it's not correct to say that they reflect Cardinal Health data. They do not. They reflect plaintiffs' expert's analysis of

DEA data.

So a factual deposition of a Cardinal Health witness on these charts, in addition to the fact that it's entirely out of time under the scheduling order in the case, would be entirely unproductive. There is no Cardinal Health fact witness who could testify from personal knowledge about what DEA did with the data or what plaintiffs' expert did with the data to generate the numbers that appears on the charts.

I would also note, Your Honor, that the summaries that have been presented with plaintiffs' motion actually were not presented in Dr. McCann's expert report. What plaintiffs have presented here is an analysis or a calculation by Dr. McCann that is nationwide. It's not limited to Cabell Huntington or to any other geographic region.

Cardinal Health was not requested to produce nationwide distribution data in this case. And Cardinal Health did not produce nationwide distribution data in this case.

So to the extent that Mr. Fuller is talking about a 30(b)(6) deposition of Cardinal Health where the witness would testify about Cardinal Health's internal distribution numbers, that would be new discovery that we would be taking in this case two and a half weeks before trial. That hasn't been requested from Cardinal Health. It hasn't been produced by Cardinal Health.

As to Dr. McCann's nationwide analysis, neither

Cardinal Health nor its litigation experts have undertaken

to evaluate all of the billions of data points that were

added by DEA before it produced the data in the litigation,

nor the tens of millions of changes that Dr. McCann made as

part of his litigation analysis.

The fact that the defendants didn't file a *Daubert* motion as to Dr. McCann's ARCOS analysis and the fact that Cardinal Health's experts didn't do a competing calculation doesn't mean that we can be forced to agree that Dr. McCann's analysis is correct.

As the Court noted at the outset, respectfully we submit that there is nothing in the Federal Rules of Civil Procedure or Federal Rules of Evidence or the stipulation that would allow the Court to, to require that from Cardinal Health.

The stipulation that Cardinal Health entered into with plaintiffs pre-trial is to allow it to work with plaintiffs' counsel to cure defects with respect to foundation of Cardinal Health produced documents.

And I appreciate Mr. Fuller saying that we've been working together cooperatively on that, and I, I share that assessment. I think we have and we've resolved many, many issues and we continue to work through those issues, but this, respectfully, is not one of them. This is not

Cardinal Health data. This is plaintiffs' expert opinion testimony, and Cardinal Health can't be compelled to stipulate to it.

What will happen if the Court denies this motion is what happens with expert analysis in every case in every Federal Court across the country. The plaintiffs will present their expert, Dr. McCann, for testimony. They will qualify him as an expert in the subject matter. And assuming they're able to do so, he will testify to his analysis.

The defendants will cross-examine him about that analysis to whatever degree they wish to do so. And the fact finder -- here Your Honor - ultimately will make a determination about what parts of the analysis are relevant and whether it accepts or rejects the analysis.

And the last point I would make, Your Honor, is that to the extent -- the Court has probably seen that the parties have now stipulated that if plaintiffs prefer to do so, they can bifurcate Dr. McCann's testimony.

The defendants have agreed that plaintiffs could call Dr. McCann once just for the purpose of introducing whatever ARCOS analysis the Court ultimately permits.

So to the extent that plaintiffs are using this motion as a method of case structure and where, for example, they want certain members in the case early instead of waiting,

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first of all, they always have the opportunity to do that
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2
     through deciding what order they wanted to call their own
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    witnesses in. But now they have the opportunity to do it by
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     agreement with the defendants exactly the way they want even
5
     if that requires Dr. McCann to testify at trial.
 6
          So the request that plaintiffs have made has no basis
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     in the rules, has no basis in the stipulations, and should
8
    be denied by the Court.
9
          I'm happy to address any additional questions that the
     Court may have, but that's all I have this morning.
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11
               THE COURT: Thank you, Ms. Wicht.
12
          Mr. Fuller, do you want to respond to any of that?
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               MR. FULLER: Sure, just briefly, Judge.
14
          As I stated earlier, we're not asking for the Court to
15
     require them to stipulate. We're simply asking for them to
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     tell us, "Hey, you're saying we have the numbers wrong."
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         And this isn't -- Judge, this isn't some complicated
18
    math. This is counting the number of pills that you shipped
19
     into each state. It's very simplistic.
20
          A 30(b) designee wouldn't be asked, "Hey, look at these
21
     numbers." The 30(b) would be asked, "What are your
22
    numbers?"
23
          And they would espouse whatever distributions Cardinal
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did of hydrocodone and oxycodone into each state, or the defendants could simply write it on a piece of paper and

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send it over to us. "Hey, these are what we think the
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2
    numbers are. These are what we believe our distribution
    numbers are."
 3
 4
          They have the data. It's their information. They say
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    using ARCOS isn't theirs, but that's where ARCOS comes from,
 6
     Judge. It comes from each of these defendants.
7
         Unless the Court has any other questions, --
               THE COURT: I don't. I realize that this needs to
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9
    be decided expeditiously. I'm not going to rule on it from
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     the bench, but I'll get an order out right away deciding it.
11
         Let's move on to the next motion. It is one of the
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     four parts of the motion in limine. It's ECF Number 1067 in
13
     the file. And the matter before the Court is part three, I
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    believe, of that motion which goes to the issue of the
15
    geographic scope.
16
         Mr. Farrell, you're going to take the lead here I
17
    believe.
18
              MR. FARRELL: Judge, I would be honored to do so,
19
    but it's the defendants' motion in limine. I'll respond to
20
     it.
21
               THE COURT: I'm reading -- your name is at the top
22
    of the script, Mr. Farrell, and I just read it off. I'm
23
     sorry. I think I've accused you before of being a defense
24
     counsel in this case.
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25 Ms. Hardin is the --

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MS. HARDIN: Yes, sir. Can you see me and hear
1
 2
    me, Your Honor?
 3
               THE COURT: Yes, I can. I can see you well and
 4
    hear you loud and clear.
 5
              MS. HARDIN: Thank you.
 6
               THE COURT: You may proceed.
 7
              MS. HARDIN: Ashley Hardin from Williams &
8
    Connolly on behalf of the distributor defendants, Your
9
    Honor.
10
          We selected this motion out of all the other motions in
11
     limine because we believe that this motion in particular
12
    will have significant impact on the scope of the evidence
13
     that is presented at the trial. I think the previous
14
     argument that you just heard demonstrates that.
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          And we expect this issue to come up as early as the
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     opening statements. So we appreciate the opportunity to
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     address this this morning, Your Honor, and hope that we can
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     receive a pre-trial ruling on this one so that your guidance
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     can inform the parties' final preparations for trial in the
20
     next few weeks and set the parameters for what will and will
2.1
    not be permissible during the opening statements.
22
               THE COURT: All right. Well, my big question to
23
    you is why shouldn't I just follow Judge Polster on this?
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               MS. HARDIN: Two reasons, Your Honor.
25
          Number one, Judge Polster's reasonings are not law of
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the case. The plaintiffs have already tried to get you to adopt that reasoning. They did so at the February argument on their motion for summary judgment on the Controlled Substances Act. Your Honor rejected that premise when he denied that, the plaintiffs' motion.

This motion has to do with what is the evidentiary nexus to Cabell and Huntington. That was never before Judge Polster. Judge Polster was not able to issue a ruling on what evidence is connected to Cabell Huntington. So that's the first reason.

And the second reason is that Judge Polster himself has revisited that ruling in the Track 3 cases which involve only the pharmacy defendants. This motion was renewed, and he clarified that what he meant by his initial ruling was that perhaps some aggregate data on the whole could be admissible without relating specifically to the Track 1 jurisdiction in Ohio. But where specific evidence is concerned, there must be a geographic and causal -- or at least evidentiary nexus between the evidence and the jurisdiction.

And here that is completely lacking, Your Honor. There is not going to be a single witness, fact or expert, who will testify, nor will a single piece of documentary evidence in the case establish that a single pill distributed by these three distributors to pharmacies

located outside Cabell and Huntington ever found their way into Cabell Huntington and caused harm there.

Plaintiffs have a few theories as to why this evidence is relevant, but none of them have any merit.

The first is their migration theory. And that is the contention that pills distributed elsewhere to places like Florida or Ohio or Kentucky or even into other places in West Virginia were trafficked into Cabell and Huntington and contributed to the public nuisance that plaintiffs allege is happening here. But that migration theory is not connected to the defendants in any way.

First of all, as I said, there's no evidence, witness or documentary, that establishes that any pills that may have been trafficked into Cabell Huntington were ever distributed by these three distributors.

McKesson, Cardinal, and ABDC are not the only distributors in West Virginia, are not the only distributors in Florida, or in any other of the states that plaintiffs have argued were the source of pills coming into Cabell and Huntington, nor is there any evidence that these defendants themselves participated in any, quote/unquote, migration.

So what migration means is criminal trafficking. It means criminal diverters went to places outside of this jurisdiction and brought pills back into the jurisdiction for the purposes of illicit drug trafficking.

There is no allegation, much less evidence, in this case that these three distributors ever distributed their products to anyone other than DEA registered and state licensed pharmacies.

And the plaintiffs' expert, their DEA expert, James Rafalski, in his report is very clear about what he thinks the source of this, quote/unquote, migration along what he terms the Blue Highway or the Oxy Express from Florida to West Virginia and that is criminal diversion. There's no allegation that the defendants themselves participated in that.

And even if there were some linkage between our pills distributed somewhere else and those that found their way into Cabell Huntington, which there isn't, but even if there were, that is not evidence that the shipments themselves by distributors into these other locales were themselves wrongful or excessive because, of course, the mere fact that a criminal traffics in drugs does not indicate that there is any wrongdoing on the part of the defendants.

What the plaintiffs have, and all that they have, is evidence that the defendants were aware of migration. But, of course, being aware that criminal activity takes place is not a sufficient evidentiary hook to admit evidence of our distributions to other pharmacies.

The very documents that the plaintiffs cite make very

clear that we're not the only entities that had that awareness; that the DEA itself was aware of migration and, in fact, Cabell and Huntington local law enforcement were aware of that migration.

So the mere fact of that criminal activity, of course, does not indicate any wrongdoing on the part of distributors. So their migration theory doesn't get them there, Your Honor.

And then I think we just heard from Mr. Fuller that they want to use certain national trends and national distribution data to serve as perhaps a benchmark for what the distributors should have known was reasonable or appropriate shipments into Cabell Huntington itself.

But that is a nonstarter and it's a reason why the argument about whether or not we have stipulated, produced our national distribution data is really beside the point because that national trend data cannot tell us anything about what was reasonable and appropriate in Cabell Huntington.

The mere -- the entire concept of an average, of course, is that there were distributions to some pharmacies that were higher and distributions to some pharmacies that were lower.

So the mere fact that someone at these three distributors might have known that the distributions to

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either a single pharmacy in Cabell Huntington or to the
region as a whole were higher or lower than some other place
in the country or some other place, frankly, located within
West Virginia is irrelevant. That's not notice that
something was amiss in Cabell Huntington because there's a
reason why we evaluate suspicious orders on an
order-by-order pharmacy-by-pharmacy basis.
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And that is because what makes an order reasonable or of typical or usual size, frequency, or duration depends on the location in which the pharmacy is located. What is it near? What population does it serve?

And, so, making those evaluations with respect to Cabell and Huntington are independent of and unrelated to making those determinations for Texas or Florida.

THE COURT: Why doesn't this come in under 404(b)?

MS. HARDIN: It is actually prohibited under

404(b), Your Honor, because there is no causal nexus. There is no evidentiary basis.

The only possible purpose that the plaintiffs can have for utilizing this information, other than the national trends that Mr. Fuller said, is as propensity evidence. And they really make no bones about this. At Pages 14 and 15 of their opposition, Your Honor, they indicate that they very much want to make an "if it happened there, it must be happening here" argument.

They want to argue that if we shipped too many pills to Florida, for example, under our national policies and procedures, then we must have also shipped too many pills to Cabell and Huntington. That is a -- that's classic propensity evidence, prior bad acts evidence that is prohibited by 404(b).

There are exceptions, of course. Plaintiffs have not argued for the applicability of any of the exceptions of 404(b). At most, what they muster in their opposition is to say that some exception could become apparent, might become available to them at the trial depending on how the evidence comes in. But that's insufficient at this point, Your Honor.

We made the motion. We, we brought forward the point that there was no evidence linking these shipments outside of the jurisdiction to Cabell Huntington and that it's prohibited under 404(b). And plaintiffs have not come forward to tell Your Honor what exception would apply or why it would apply.

So 404(b) is prohibitive of this evidence. So it's irrelevant. It's prohibited under 404(b). It's actually going to prejudice the defendants. If, if the plaintiffs are allowed in this case to base a liability finding for nuisance in Cabell Huntington based on conduct that we took in other places, then we're going to be subject to repeated

verdicts for the same conduct because those other jurisdictions have also sued us, Your Honor.

The State of Florida has sued these three distributors. And, so, we're going to fall to account for our conduct, our shipments into Florida. And if there is evidence that they were wrongful or in excess of what was allowable, the State of Florida is going to litigate those claims.

As you know, 54 out of 55 West Virginia counties have sued these three distributors. And, so, if we've made inappropriate shipments to other places, that's going to be litigated before the Mass Litigation Panel.

And, so, without a connection between conduct in those jurisdictions and this jurisdiction, these plaintiffs cannot hold us liable for that same conduct without subjecting us to repeated judgments, and that's inappropriate.

And, of course, Your Honor it's going to be an enormous waste of trial time. We're already slated to be together for 12 weeks just talking about the issues that are relevant to Cabell and Huntington. And if we have to devolve into a bunch of mini trials talking about the various shipments that were made to Florida or Ohio or Kentucky, we're going to be here for a long time because defendants do not agree that any of our conduct was wrongful in any of those places anymore than we agree that it was wrongful in Cabell Huntington.

And, so, if plaintiffs put forward evidence of shipments to other places, defendants are going to want to defend themselves. And we will be required to defend ourselves and admit evidence about those shipments and it's going to take trial time that we don't have and it's irrelevant. And for that reason, we ask that Your Honor exclude evidence of our out-of-jurisdiction shipments.

THE COURT: Thank you, Ms. Hardin.

Mr. Farrell, I now have you appropriately aligned in this case with the defendants no longer and you may respond.

MR. FARRELL: Thank you, Your Honor. I will note that there are other Farrells that practice law in this state that would probably not mind being aligned on the defense side.

Thank you, Judge. This is an important issue. And at the outset, I wanted just to affirm that, yes, the geographic scope has been the subject of not only discovery, but evidentiary rulings in other courts.

But before we kind of get there, I want to -- I don't want to engage in closing arguments right now. And, so, some of what defense is raising is the weight of evidence and the, the probative value of some of this evidence.

And I'm going to try to stay away from that as best as
I can and point out the irony initially that the defendants
are filing a motion in front of you to prevent you from

hearing evidence as a fact finder. And it seems like the cat's out of the bag because you're the jury and the Judge.

That being said, the Fourth Circuit has already put together some of these rules in saying that, you know, prejudice for a bench trial, you get to weigh the evidence.

And, so, I'll say this at the outset. I am acutely aware that the plaintiffs are going to need to follow the rules of evidence. We're going to have to lay proper foundations. We're going to have to be methodical. We're going to have to stay within the facts and present to you a case in order for us to prevail.

And, so, in general, what I can tell you is this. The reason this information is important is because all three defendants have acknowledged that they have a duty to identify and monitor suspicious orders. All three defendants have distribution centers across the country. And all three defendants have acknowledged on the record that their Suspicious Order Monitoring System, SOMS for short, is systemic.

So for whatever successes these companies have in complying with their duty, it's as a result of a systemic process. And whatever failures it has is the result of a systemic process.

So we intend to present evidence that what happened with the volume of pills in Huntington, Cabell County was

not a mistake. It was not an isolated event. It was the result of a systemic failure by each of the three defendants to comply with their regulatory duties to identify, block, and report suspicious orders.

Suspicious orders are defined in the Code of Federal Regulations as orders that are of unusual size. In order for us to determine what is unusual, we're going to have to define what is usual. And to do so, what we have done is we have hired experts to come in and review this data and present it in a logical, methodical way to you so that we can identify the facts of national, regional, state, and local patterns.

And I will point this one thing out. The defense is arguing that the national trends, the benchmark, cannot tell us anything about what is suspicious. And I will point out that that very argument is the basis of the DEA investigating them, sanctioning them, and fining them for the past decade.

I'll also point out that the defendants concede that they were aware of migration. And I will point out that there is a case that we will be discussing at great length called *Direct Sales*, which is a case that was actually provided to the defendants by the DEA. It's dated 1947 I believe. And it talked very much about this awareness and the culpable liability that comes with such awareness.

So, yes, we intend to discuss migration. We understand we have an evidentiary burden. We will put on evidence through our own internal documents. We will put on evidence of the Oxy Express and Blue Highway through the DEA documents. And we're going to put on evidence of the Oxy Express and Blue Highway through the defendants' own internal documents.

We're going to put on evidence that the problems that were -- that, that resulted, the consequences and carnage that happened in Cabell Huntington is the result of a systemic failure.

We'll be able to establish that the volume of pills in Huntington Cabell weren't being diffused by going in other places because our surrounding counties were immersed in pills themselves.

So, in general, an argument under 404(b), we understand that, yes, we have obligations under the rules of evidence regarding relevance and prejudice. You as the finder of fact will be the gatekeeper and I'm sure will be very clear as to when you've heard enough. And you'll be able to give appropriate weight to the evidence.

THE COURT: And you're relying solely on 404(b) to get this in, Mr. Farrell?

MR. FARRELL: Well, no, we think this is direct evidence of their systemic failures. In addition to 404(b),

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we'll be able to show that this was not a mistake, that this
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    was part of their plan.
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               THE COURT: When Judge Polster ruled it was
     admissible, did he rely on 404(b) if you know?
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              MR. FARRELL: I do know. His direct quote I
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    believe -- I'm looking at Page 15 of our response, the end
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    of Page 14 bleeding onto Page 15. And what he basically
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     says is that evidence outside of Track 1, CT1, comes in.
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     says this is particularly true because there is evidence
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     that defendants acted pursuant to practices and policies
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     that were national in scope.
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          So I don't have the order in front of me but, in
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     general, there not only is direct evidence, but we're also
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    going to be able to show the other counties and cities and
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     states for purposes of benchmarks. So I don't think this is
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     a back door attempt, Judge. This is blunt, direct evidence
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    of our case in chief.
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               THE COURT: Did Judge Polster revisit this issue
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     in the Track 3 cases?
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               MR. FARRELL: Now, my focus -- as you know, I am
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     the co-lead of the national litigation, so I'm going to
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     feign a little bit of, of a lack of knowledge on this.
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          I believe what is happening in CT3 is that we're still
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     in the discovery phase, and I fully anticipate that this
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     issue will be re-litigated. But the Judge entered a 75-page
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evidentiary order that makes it pretty clear that he's going
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    to allow this into a jury.
          And all I'm suggesting is that if we get to trial and I
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    have failed to put on proper evidence, I'm sure the defense
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    will be keen to object and you'll be, you'll be prompt in
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     sustaining the objection.
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               THE COURT: Well, I've got Judge Polster's order
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     in my hand and he very emphatically on the last page
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     indicated that in his view, the rulings he made should apply
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    to all the remanded cases.
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          I realize that's not binding upon me, but I have a lot
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    of respect for the work that he's put into it and I just --
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     I'll be asking your opponent this. But shouldn't I just
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     follow Judge Polster?
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               MR. FARRELL: Well, Judge, I --
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               MS. HARDIN: Your Honor -- I'm sorry. Did you
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     want Mr. Farrell to address that or me?
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               THE COURT: Yeah. I'll give you an opportunity in
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     a minute, Ms. Hardin.
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          Go ahead, Mr. Farrell.
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               MR. FARRELL: Yes.
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               THE COURT: Your answer was probably "yes."
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    Right?
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               MR. FARRELL: Yes, I would like for you to follow
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     Judge Polster's orders to the extent they help my case.
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I would ask you not to follow his orders to the extent that it hurts my case.
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THE COURT: Fair enough.

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Ms. Hardin, do you want to respond to that?

And if you're not through, I'll get back to you, Mr. Farrell.

Go ahead, Ms. Hardin. The issue is whether I should follow Judge Polster or not.

MS. HARDIN: You're certainly not required to follow Judge Polster. I think Your Honor just recognized that. His rulings are not law of the case. As a general matter, we've dealt with that already. We've put that issue to the side. I take it from your rulings already you don't adhere to the notion that they are law of the case.

But particularly when we are talking about motions and what evidence has been adduced in this case, there's no reason to follow Judge Polster. He didn't have this record before him. He had a different record before him with different evidence and different facts.

And, so, we ask Your Honor to make your own rulings based on what the evidence is here.

But to Your Honor's question has he reconsidered or re-evaluated, the answer is, yes. The Track 1 ruling is not his final word on this issue.

On November the 3rd of 2020 -- and we cite this at Page

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12 of our reply which is in this case Docket 1154 -- he says
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     this. He said while he would permit broad-scope aggregate
    evidence related to things like nationwide trends and
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     shipments, which we certainly disagree with but that's his
     ruling, he said extraterritorial evidence must have some
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     demonstrable nexus to the plaintiffs such as the fact that a
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    distribution center located outside of the plaintiff
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     counties distributes prescription opioids into them.
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          So even Judge Polster -- again, we disagree with the
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    broad scope of his ruling. But even he recognizes that when
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    you're talking about specific evidence, it needs to have a
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     demonstrable nexus.
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          I have other points to respond to Mr. Farrell, but I'm
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     not sure if he was finished or if he's got more to say or I
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     should continue.
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               THE COURT: Okay.
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          Do you have more to say, Mr. Farrell?
               MR. FARRELL: I think I made clear I have more to
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          But right now, Judge, I think I've said what we need
     say.
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     to.
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               THE COURT: All right.
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               MS. HARDIN: Just a couple of points then if I
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    might, --
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               THE COURT: Yes, sure.
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               MS. HARDIN: -- Your Honor.
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On the systemic failures point, if, if we have national policies and procedures and they impacted what happened in Cabell and Huntington, this motion is not trying to prevent the plaintiffs from talking about that evidence.

If our Suspicious Order Monitoring System on the whole, as they were implemented in Cabell Huntington, had certain failures or inadequacies, I feel certain that Mr. Farrell and his team are going to try and point that out and this motion isn't going to try to keep them from doing so.

But what pills we shipped to other places pursuant to those policies just is not relevant. It doesn't matter if we shipped too many, too few, or just the right amount of pills to a pharmacy in Florida or Ohio or Kentucky. That does not mean that we shipped too many or too few or just the right amount to Cabell and Huntington.

And, so, if they want to talk about why our policies and Suspicious Order Monitoring Systems were insufficient with regard to the decisions we made with respect to Cabell Huntington pharmacies, they may do so. But they don't need to tell you how we operated under those policies in some other place. That's not going to inform what we did here.

And we've not put mistake at issue, Your Honor. We've not argued that our distributions into Cabell Huntington were a mistake. So they don't need to introduce evidence to show that we didn't make a mistake. We're going to defend

1 our Suspicious Order Monitoring Programs on their merits.

2 But we have not said that somehow mistake or identity or

3 opportunity is at issue. So it just doesn't pull in 404(b).

And, again, the notion that unusual size, frequency, or duration is (video inaudible) for a Cabell Huntington pharmacy based on what might be happening in Florida or Texas or New York City is incorrect. That's not how suspicious orders are evaluated for the reasons that I

already said. They are locale and pharmacy specific.

And to the issue that this is an issue of weight of the evidence, there is no evidence. We haven't talked about any evidence that connects the distributors to the migration theory. They have evidence of the migration theory to be sure, but they don't have anything that connects us to that theory. So that is not a sufficient basis on which to admit our distributions from outside of the jurisdiction.

Thank you, Your Honor.

THE COURT: All right. I'm going to take this one under advisement. And let's go on to the last thing on the, on the agenda here. That's the defendants' motion to exclude the marketing opinions of four experts. And Kim Watterson is going to present this, I believe, for the defendants.

MS. WATTERSON: That's correct, Your Honor. I will check. Am I transmitting both audio and video to you?

1 THE COURT: You are, Ms. Watterson.

MS. WATTERSON: All right. Well, thank you. Kim Watterson, Reed Smith. I'm here arguing today, though, on behalf of all of the distributor defendants in this case.

And I think like the geographic scope issue that was just argued, we really appreciate you hearing from us on marketing because it too will help shape the scope of the evidence in the case, or your rulings will, and it will have an impact perhaps on the length of the trial.

Your Honor, it always helps me especially with a motion like this where there's several experts implicated, several moving parts in terms of the number of different *Daubert* criteria at play to set forth a little bit of an inventory before I start.

So there's going to be three basic parts to my argument today.

First, I'm going to explain why three of plaintiffs' marketing experts -- and I'm going to shorthand for now and call them marketing experts - Keyes, Lembke, and Kolodny - are not qualified to give an opinion on what again I'm short-handing as marketing for now.

Second, I'm going to explain why those three experts, Keyes, Lembke, and Kolodny, as well as plaintiffs' other marketing expert, Jakki Mohr, on these experts plaintiffs have not satisfied their burden. And, again, it's

plaintiffs' burden under *Daubert* to demonstrate reliability and relevance when it comes to all four experts.

And, third, I'm going to talk about Professor Mohr in and of herself and explain why, given her own descriptions of the limitations in her opinion and what remains after we look at those limitations, the testimony that she's going to provide is simply not helpful to the Court. It doesn't meet the criteria that expert testimony must be helpful to the finder of fact. And, of course, Professor Mohr's testimony doesn't fit the circumstances of this case.

And when we talk about Professor Mohr, I want to foreshadow this. This Court's recent April 8th order is going to come into play in that analysis.

A little bit of context before we begin to talk about these experts.

Throughout this opioid litigation, plaintiffs' marketing liability theory has focused on manufacturers.

That's reflected in the complaint that they filed in this case.

If you look at Paragraphs 372 through 669, there's a discussion of manufacturers' alleged marketing activities. Their experts in this case until recently, that is, their marketing experts have focused on manufacturers' conduct.

But now that we find ourselves in this case, and a few others, where the distributors are the only defendants,

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plaintiffs are trying to read the distributors --
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               THE COURT: Is it stronger against the
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    manufacturers than it is against the distributors or is it
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     comparable?
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              MS. WATTERSON: It is -- pardon me, Your Honor?
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     Did you say is it weaker or is it comparable?
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               THE COURT: All of the above. Is it -- I'm under
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     the impression that the plaintiffs' evidence is better
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     against the manufacturers, who aren't in this case anymore,
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     than it is against the distributors. And I'm just asking
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    you if you agree with that or not.
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               MS. WATTERSON: I think it's certainly better. I
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     think manufacturers have been the focus of their marketing
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     liability theory. I will argue today they do not have
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     expert testimony on the critical link in that theory.
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          You know, the marketing liability theory is that
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     defendants engaged in marketing activities that caused
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    physicians to prescribe more opioids. You know, stated in
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     its simplest fashion, that marketing drove the supply of
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     opioids by misleading physicians into prescribing more
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     opioids, not paying attention to the risks of opioids,
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    marketing that overstated the benefits.
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          So that's the marketing liability theory which, again,
     I don't want to get caught up on this distinction between
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    manufacturers and distributors by way of setting forth the
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context, but I do think it is an important context. And I also think it lays the groundwork for why their marketing experts, and particularly their marketing causation experts, really don't have anything to say that passes *Daubert* muster when it comes to distributors.

So let me start with qualifications. You know, we talked a little bit about Judge Polster's order. Your Honor, you asked us in our papers to indicate whether Judge Polster has ruled on a particular issue that's before you. So we did that.

Judge Polster ruled as to two of these experts, Lembke and Keyes, and the reasoning applies equally to Kolodny, that they do not have the requisite experience, education, and training to render an opinion on marketing and its effects on prescription prescribing.

Now, we're not just talking about Judge Polster's opinions because we think it's the law of the case because we don't think Judge Polster's opinions are law of the case. But Judge Polster was perfectly correct, and we independently demonstrate to you in our papers that Lembke, Keyes, and Kolodny do not have the requisite experience.

Lembke is a medical doctor. She deals with addiction and pain. She offers some opinions in this case that do not deal with marketing and are not the subject of this motion. But she has no background whatsoever in marketing.

Plaintiffs point to a book that she authored. But authoring a book that mentions marketing does not qualify her as an expert under the *Daubert* criteria. Plaintiffs say repeatedly that Lembke, as well as Keyes, and as well as Kolodny, are reading up on things in order to prepare to testify in this case.

Well, this Court's opinion in Salazar has made very clear you don't gain expertise just by getting ready to testify in litigation as an expert. You don't become an expert outside of your regular field just by reading up on things.

Plaintiffs point to a class that Professor Lembke teaches on the opioid epidemic, as she calls it. One day she teaches a class throughout the entire course, just one day, where she touches on marketing. That doesn't make her an expert under *Daubert*.

The same goes for Keyes. Again, she's an epidemiologist. She has no credentials that qualify her to talk about marketing, and specifically whether marketing here drove prescriptions of opioids or physicians' prescriptions.

Again, what plaintiffs have to say is she's working hard to get ready to go to testify. That doesn't pass Daubert muster.

Interestingly, with respect to Keyes, this crash course

that she might be engaging in on marketing, by her own
admission she isn't even evaluating -- and we'll talk more
about this when we get to the reliability criteria. But she
doesn't even -- she disclaims having any knowledge of
distributor marketing. And she hasn't evaluated the
specific marketing materials that, that are at issue in this
case.

And, again, the same goes with Dr. Kolodny. Again, he is a physician. He treats OUD. He is not positioned to testify on marketing and marketing causation.

Plaintiffs may raise, because we're talking about other judges' opinions, that the judge in Oklahoma issued a one-liner allowing Dr. Kolodny to testify. But there's no reasoning in that opinion, and nothing that suggests that he's qualified under *Daubert*.

And, certainly, that pales in comparison to what I will say is Judge Polster's well-reasoned analysis here and what we argue in our papers looking specifically at what these folks have by way of background, what they're lacking when it comes to expertise in this area, and why under the controlling case law they simply don't meet the qualification criteria.

Your Honor, I'm going to move now on to the reliability prong. Maybe fit and reliability are the two most important touchstones under *Daubert*.

None of these experts offer a reliable opinion on marketing causation, again defined as, in accordance with plaintiffs' liability theory, the theory that distributors engaged in marketing that drove an increase in the supply of prescription opioids because they convinced doctors through misleading statements to prescribe more opioids to more patients in higher doses and for a greater multitude of conditions.

So, again, that's the focus. That's the frame. That's what these experts need to offer a reliable opinion on based on a methodology, based on data, not simply speculation and surmise.

So for starters, let me point out that Jakki Mohr, the only expert of the four who has any background in training, in marketing, she has essentially disclaimed having any opinion on causation. She said her opinion is limited to whether distributors engaged in marketing activity; were these activities marketing; can they be considered marketing.

She has no opinion that any of these activities were false or misleading. And she has no opinion on whether any of this marketing had an impact on the overall levels of opioids prescribed, and certainly not on the overall level of opioids prescribed in Cabell and Huntington.

Those experts that do purport to have an opinion on

marketing causation - Keyes, Lembke, and Kolodny - they didn't employ any methodology at all. They didn't talk to any physicians anywhere, and certainly not in Cabell and Huntington, -- and we'll talk about that later specifically with respect to the fit criteria -- to determine whether and to what extent any of these marketing activities, whatever they may be, whether and to what extent they influenced prescribing. They just didn't do it.

You know, there isn't any pile of data that they looked at to say, "Hey, physicians, did you get this information? Did this move the needle?" There is no underlying data set, and there was no methodology.

It's just, well, you know, if folks engage in marketing and tout the benefits of opioids and underplay the risks which, of course, is plaintiffs' theory, of course that impacted physicians.

But I want to draw this Court's attention to the case of In Re: Actos and In Re: Diet Drugs, as well as this Court's opinion in Salazar as well. It says it's really just speculation, you know. This is In Re: Actos. They excluded marketing causation -- and I'm going to read what it says -- because there was no foundation laid to show that the experts knew what a physician actually knew, what marketing materials the physician actually received, and whether and to what extent the physician was impacted by

those marketing materials.

So there, there is no methodology at all. It really amounts to an *ipse dixit*. They admit, and we cite in our brief, all of the pages of the depositions where each of these experts admitted they don't know whether any distributors distributed marketing material. They don't know whether any doctors saw such marketing material. They don't know whether doctors relied on or the extent to which they relied on these marketing materials in making prescribing decisions.

Now, plaintiffs come back and say, well, you know, we don't have to tell you which doctors. We don't have to identify specific doctors that relied on these materials.

Your Honor, that's not our point. We're not faulting the experts because they don't give us an inventory or list or count up the numbers of doctors who say they relied on these marketing materials.

It's simply that they don't have any methodology at all. There wasn't any examination of underlying facts from which they could draw their opinions. So it's *ipse dixit*. It's guess. It's surmise. It's, it's speculation that somehow these materials must have moved the needle.

That's not what *Daubert* requires. There has to be a methodology. And there really is none here when it comes to the question of whether they can opine that distributor

marketing moved the needle and influenced physician prescribing.

Another thing, Your Honor, that's missing from the Keyes, Kolodny, and Lembke opinions — and it's nothing, by the way, that Professor Mohr admits that she didn't do. You know, all of these experts admit and agree that there are other factors that influence prescription — physician prescribing; manufacturer conduct, DEA, FDA, physicians' own judgments, et cetera, et cetera. So all of them agree that there are other drivers of prescription prescribing.

When asked whether they did any assessment at all, well, what role does this distributor marketing, if any, play in the equation, none of them evaluated that. Mohr said she didn't do it because she said that wasn't her task. And she said in order to reach a conclusion on that question, you would have to do, and her quote was an economic, an econometric study that would take some time to sort out. "That wasn't the scope of my task."

Well, Keyes, Kolodny, and Lembke didn't do that either. So, again, it's *ipse dixit*. I hate to keep using that word out of the Supreme Court precedent, but that's what it is. They say we understand they did some marketing. It must have moved the needle. But we can't tell you how much and we can't tell you based upon a methodology and the examination of data, a close examination of the marketing

material, and talking to physicians that it moved the needle. We just suspect and surmise, well, it must have.

Now, also missing from the marketing experts' opinions is fit. All of them admitted that they didn't do any look into Cabell County. They don't know whether any of the so-called marketing materials and marketing activities -- and, Your Honor, you'll notice I'm calling them so-called because for the purposes of this motion now, we're not probing into those materials and deciding, you know, whether or not, or arguing that they weren't disseminated, whether they were disseminated, and whether they really amount to marketing.

But whatever the so-called marketing might be, none of the experts know whether any of these marketing materials made their way into Cabell and Huntington, whether physicians in Cabell and Huntington looked at them, whether they resulted in a prescription being written for someone in Cabell and Huntington. They were all very honest about it. They said that they didn't look into this jurisdiction.

So what they have to say simply doesn't fit this case brought by Cabell and Huntington looking at impact in this jurisdiction. So they all flunk the *Daubert* test when it comes to both reliability and fit.

Now, I want to talk a little bit about Jakki Mohr because interestingly, Your Honor, I mentioned to you that

she disclaims having any opinion at all on this marketing causation question. She says, "All I'm really doing is answering the question of whether distributors engaged in marketing."

She's not opining on whether any of the so-called marketing was false. And she says she didn't do any analysis to determine whether or not that marketing, in fact, moved the needle and led to physicians overprescribing.

So all we are left with when it comes to what Jakki
Mohr has to say is an inventory of activities,
communications, programs that she looks at and she says,
"Yes, that amounts to marketing," not that this marketing
did anything to cause prescribing, not that it's false.
That meets the definition. "I'm a professor of marketing.
This meets the definition of marketing and this is how
marketing works."

Helpfulness to the trier of fact, as this Court has said in its own opinions, is the, is the touchstone of the Rule 702 analysis. How is that helpful to the Court? What remains is she's going to get on the stand and she's going to be the vehicle through which plaintiffs put on an inventory of activities and communications. And then she's going to say, "In my expert opinion, I think that's marketing."

Well, first of all, whether an expert says it's marketing or not doesn't inform the central question here.

The central question here is whether these communications or programs, whatever you call them, marketing or something else, influenced prescribing habits such that there was a greater supply of opioids in Cabell and Huntington. She can't say that.

And as this Court noted in its April 8th opinion, experts -- expert testimony isn't the vehicle for a party, here in this case the plaintiff, to get into evidence facts about corporate documents or corporate activity.

So at the end of the day, that's really all that we're left with when it comes to Jakki Mohr. And she should, therefore, be excluded in her entirety. She has nothing to say about causation. She has nothing to say about falsity. She doesn't fit. She has nothing to say about Cabell.

And what's left of her testimony, again, this inventory of activities and then affixing the marketing label on them, that's not helpful to the trier of fact. That's not going to help Your Honor determine whether plaintiffs' marketing liability theory carries the day. And it's also not the proper vehicle to put in the evidence about these so-called marketing programs.

I think, Your Honor, I will stop there -- there's a lot more to say, there's a lot going on with marketing -- and

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I'll answer any questions that you might have.
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               THE COURT: I'm going to hear from Ms. Bierstein.
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          Thank you, Ms. Watterson.
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              MS. BIERSTEIN: Sorry, Your Honor. I had to
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    unmute.
              I had to remember to unmute. But can you hear me
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    now?
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              THE COURT: I can hear you loud and clear, Ms.
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    Bierstein.
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              MS. BIERSTEIN: Okay, great. Thank you. Good
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    morning, Your Honor. Andrea Bierstein, Simmons, Hanly,
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     Conroy, arguing for the plaintiffs.
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          This motion to exclude the marketing evidence of
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    plaintiffs' expert, as I'm going to try to demonstrate to
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    Your Honor, should be denied because each of the experts at
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     issue --
               THE COURT: Let me interrupt you. Judge Polster
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     ruled at least two of these experts out. How -- why
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     shouldn't I do the same thing?
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              MS. BIERSTEIN: Well, I was going to get to Judge
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     Polster, but I'll get to him first since that's the focus of
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    Your Honor's question.
          I noticed that all counsel, Ms. Watterson, Mr. Farrell,
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    we all seem to agree that you should follow Judge Polster
    when we like what he does and that you shouldn't follow him
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when we don't like what he does.

But, you know, looking more specifically here, I want to say a few things about Judge Polster's ruling.

First of all, Judge Polster's ruling was extremely narrow. The relief the defendants are asking for here is quite a bit broader.

What Judge Polster did was to exclude very limited amounts of testimony. And you see in his opinion it's like, "Well, I'm not letting you say this paragraph and this paragraph." It's very surgical.

And a lot of what's at issue here is simply either not the subject of his ruling -- certainly was not -- the breadth of what they're seeking to exclude is, is much broader than what Judge Polster addressed.

But even as to what Judge Polster did address, the record before him was quite different I think in, in certain important ways from the record before Your Honor. And I know this is kind of going a little bit backwards in terms of, you know, the way I was going to present this.

But I want to start with what Judge Polster said about Dr. Keyes in particular because one of -- because Judge Polster's main criticism of Dr. Keyes -- and this is quoting from his opinion -- was that her report, he said, does not indicate that in formulating this opinion, Keyes performed the methodology that is standard in the scientific process of her field of expertise.

In other words, Keyes has not shown that she applied epidemiological methods to determine that a cause/effect relationship may be inferred from the study she cites.

So he didn't say -- I want to say he didn't even say she hadn't used the methods, but he said her report didn't show that she had.

And the report here is different because if you look on Pages 29 to 31 of Dr. Keyes' report in this case, different from her report in CT1, you will see that Dr. Keyes isn't just reporting on articles, and I'm going to talk some more later about this business of reading up, that she's not only -- she's analyzing the literature specifically using the epidemiological tools to do that. She's assessing the extent to which the studies at issue are epidemiologically sound.

And, so, what -- so what she's doing here is what she didn't do in front of Judge Polster. She's demonstrating that she's using her epidemiological expertise and her epidemiological methods in looking at epidemiological literature in order to reach her conclusions.

So I think she's cured of what Judge Polster was concerned about which is, yes, what she's doing here is epidemiology. And she obviously had not made that clear enough in her CT1 report. We think that in the CT2 report she has demonstrated to the Court that that is what she's

doing.

And I am going to come back and talk some more about what she's doing and why it is epidemiology in just a minute. But I did want to answer Your Honor's question.

I think with Dr. Lembke the situation is a little bit different, but not that different. It's true that the record in front of Judge Polster included original research that Dr. Lembke had performed before writing her book.

By the way, you know, Ms. Watterson, the defendants keep saying nobody ever talked to any doctors. Well, Dr. Lembke did. When she researched her book, she did extensive patient and physician interviews, as well as analyses of published literature about prescribing.

And although a lot of that was in the record, Judge Polster didn't mention any of that research when he found her book to be an insufficient basis. And I don't know whether he had focused on it or whether it sort of, you know, didn't come to his attention. He doesn't say, "Well, yes, she did all this research, but it's not enough."

And I want to say -- and this is where I want to talk about reading up on something which we hear a lot about, how you can't just read up to become an expert.

Well, that's not exactly true. And the reason it's not exactly true, Your Honor, is that a literature search is a respected and well recognized methodology that has passed

Daubert on numerous occasions and it's been recognized by numerous courts.

There is a limitation on a literature search which is that you need to have the right expertise. A medical doctor can read medical literature and analyze it and figure out:

Is it sound? Was this study done correctly? Does this make sense? If I read the same medical study, I could not bring my expertise to bear to understand it.

So what you need when you're reading up on something, you need to be reading up in your own field. And as long as you're reading up in your own field so that you are -- you have the same or similar expertise to the literature that you're reading, you don't have to have had an opinion or know the specific issue beforehand. You can read the literature and, and form an opinion based on it if you bring to bear the correct expertise.

And I want to circle back to talk about why Drs.

Lembke, Kolodny, and Keyes do bring the right expertise such that the component of their work that involves literature search is, is not a problem because they have the qualifications to do it.

Before I get there, I'm going to circle back to a couple of kind of preliminary points.

I did want to talk just for a second about Rule 702 and Daubert generally. I'm sure Your Honor knows the purpose of

Daubert is for the Court to act as a gatekeeper with respect to expert testimony. And as Mr. Farrell has already alluded to, in this circumstance with a bench trial, the Court is essentially a gatekeeper for itself.

So it's not to say we don't have to meet the requirements of, of Rule 702. But as Your Honor recognized in the *Grant Thornton* case, this applies differently in a bench trial where the relevance and the helpfulness of the evidence in particular can be more easily assessed at trial in the context of the rest of the evidence that comes in.

And issues that may arguably go to weight rather than admissibility or sufficiency, some defects in the methodology or smaller gaps in qualification are also best dealt with in the context of, of the trial because it's in that context that the Court can see how helpful is it, how relevant is it, how, you know, big a problem is this, you know, methodological flaw that cross-examination has brought out.

And because the Court is the gatekeeper for itself here and not for, you know, a jury that may be swayed by the mere fact that somebody with a Ph.D. is on the stand, I think the Court needs to keep that in mind in applying this.

I also want to note that although the defendants' motion is a little bit vague about exactly what parts of these experts' opinions they're trying to include -- trying

to exclude. But the scope is pretty broad. They're asking the Court to exclude large swaths of evidence, as I already mentioned, way broader than what Judge Polster did, which was very narrow.

But I also want to mention to Your Honor that the total amount of testimony at issue here is small. This is not going to shorten the trial by substantial amounts. And that's because this motion contains no challenge to the opinions of Dr. Lembke about addiction, about pain medication, about the falsity of defendants' misrepresentations, the host of other topics in her expert report.

It includes no challenge to the rest of the opinions of Dr. Keyes about opioid use disorder, diversion, the connection between oversupply and harms. And it doesn't --well, I'll get to the issues with Dr. Mohr because they now say what I think was not clear in their motion, that they think she should be excluded in her entirety. But it doesn't address the rest of Dr. Kolodny's opinions about oversupply and the opioid epidemic.

So we're talking about a smaller amount of testimony. And it's in that circumstance again that it's especially appropriate for the Court to hear all of the evidence and determine in the context of the trial whether to consider this evidence and, if so, what weight to give it.

It would not make sense for the -- for even in a bench trial for a Court to do that for weeks and weeks of testimony that takes up everyone's time only for the Court to say at the end, "Well, I'm sorry, but really that, you know, I don't see it."

But where it's a small amount of testimony and where I think I'm, I'm hoping to demonstrate we do meet the requirements, the threshold requirements, the Court will be better positioned in the context of the trial to assess the relevance, the fit, the methodology, and all the, all the rest of that.

Before I, I talk more specifically about the qualifications of our experts, though, I do want to hit two other points.

And the first has to do with the term "marketing" which is in some ways a little bit misleading. But I think it also helps understand why Dr. Mohr's testimony is so helpful here.

When we think of marketing, I think most of us think of advertising and maybe a little bit of PR. But what we're talking about here is really much broader, just how do you change perceptions of a product so as to increase (video inaudible) it fails. How do you get the relevant decision-makers to think differently about your product.

Now, that includes advertising and branding and

promotion, but it includes a lot more. It includes issue management. It includes education. It includes all the ways that you change people's minds in order to make them receptive. It may include lobbying. It may include working with medical societies. It may include getting your message into continuing medical education.

It is a very broad -- the activities at issue here are very broad. And the word "marketing" does encompass that, but it's not necessarily what most of us think of as marketing.

And Dr. Mohr, who is an expert in exactly this issue, explains that in her report. She explains how broad marketing is and she explains the role of issue management and other aspects of marketing.

So that when she says, "I've looked at what the defendants did and, yep, that's marketing," you know, and they say, "Well, that's not helpful to the fact finder," well, it's helpful to the fact finder when the marketing strategies go way beyond what we usually think of as marketing.

It's also helpful to the fact finder because the other thing that Dr. Mohr is an expert in is something called channel management. And this has to do with the way manufacturers, distributors, and retailers work together in order to get -- to market a product, to increase sales.

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They have an aligned interest. The manufacturer wants to increase sales. The distributor makes a profit off how much is sold and so does the retailer. And the ways in which the different actors in a supply chain work together in order to do that is critical and important information that is not simply obvious, that is not known to everyone who doesn't have a marketing background. And this --THE COURT: Did Dr. Mohr's, Professor Mohr's opinion reach the issue of causation? Did she have an opinion on whether the marketing -- the distributor marketing tactics were actually a cause of the damage, damages alleged by the plaintiffs in the case? MS. BIERSTEIN: I don't believe that she does. think it is the other experts who can identify the causal connection. She's in a position to identify the scope of those activities and how those activities work in the marketplace. THE COURT: So if the other experts on this point are excluded, there would be no evidence of causation. Right? MS. BIERSTEIN: I don't think that's true, Your Honor. I think there would be no expert evidence of There's plenty of other evidence of causation. causation. For example, a lot of -- certainly, the manufacturers, and I think perhaps the distributors, and I'm going to get

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in a minute to that connection, did their own internal
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     analyses of causation.
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               THE COURT: That was a bad question. I should
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    have limited it to experts. So you don't need to go further
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    on this point --
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               MS. BIERSTEIN: Okay, all right.
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               THE COURT: -- because I asked a bad question.
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               MS. BIERSTEIN: Yeah, okay.
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          So -- but, yes, it is the other experts that we're
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     relying on to make this, this causal connection.
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          And before I turn again to why these experts are
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    qualified to do that, I do want to say that -- and I think
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     this is a really important point to understand what's going
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    on with these experts.
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          One of the defendants' points is that they say we
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    haven't distinguished and our experts haven't distinguished
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     the effect of distributor marketing from that of
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    manufacturer marketing.
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          And I want to say, Your Honor, I think that gets to the
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    heart of the issue because that criticism is really a
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    non-sequitur. It misses the point.
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          What we're saying here is that the defendants and the
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    manufacturers were acting in concert on a single marketing
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     scheme to alter the perceptions of doctors and patients
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about the risks and benefits of opioids.

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The distributors offered marketing services to the manufacturers to help the manufacturers with their marketing. It's not two separate streams of marketing where the experts need to say what was the effect of this stream, what was the effect of that stream.

THE COURT: So if you succeed on this point, you bootstrap any weakness you have in the evidence against the distributors by tying them to the marketing -- to the, to the manufacturers against whom your evidence may be stronger. Right?

MS. BIERSTEIN: I think that's right, Your Honor.

And I -- we understand -- we do still need to prove that to
you. We have to prove that they acted in concert. And we
have to prove to you that it was one scheme.

And we think the documentary evidence at trial will show that. But once that's established, that's correct.

Now we're in the world of looking at what was the effect of that scheme.

And here's the interesting thing. Your Honor refers to that evidence about the manufacturer scheme. You know, you've said it's better or stronger. But what I think is interesting about it is that there really isn't a dispute about the causal connection between that marketing activity and the huge increase in prescribing and sales of opioids that occurred in West Virginia and around the country.

Defendants have been saying all along that marketing changed the standard of care and caused an increase in prescribing. They've just been saying it was the manufacturers' marketing, not ours. But they have -- you know, when they thought we were only talking about the manufacturers, they were happy to say our SOMS failures didn't cause the oversupplies they argue. The oversupply was caused by those manufacturers and their marketing scheme.

So they've been saying for a long time that it's the marketing scheme that brought all the pills to West Virginia, not our SOMS failures. Now, we, of course, are saying it's both. It's the SOMS failures and the marketing scheme.

But, more important, we're saying you defendants, you were working with them on the marketing scheme. You were in concert. And you are just as responsible for it when you signed up and said, "Hi. How can we help you with your marketing? We'd like to be part of that." And, so, that I think is an important crux of, of this case.

Now I want to turn to what I think is the heart of defendants' motion, their attack on Drs. Lembke, Keyes, and Kolodny, which I think is the misconception that marketing professionals are the only ones who can assess the effectiveness of marketing here, that they are the only ones

who are qualified to talk about causation.

And, you know, so we have a situation. A company or a group of companies spends millions of dollars on marketing. The sales go through the roof. But the theory is that only a marketing professional can tell us if the two things are connected, if the increase -- if the marketing plan and all the money they spent actually caused the sales to go through the roof.

Now, I will say there's a superficial appeal to that argument. After all, we all know that correlation isn't necessarily causation. So, you know, you put in the money, the sales go up. How do we know? Someone has to assess whether there was a connection, whether the sales would have gone up anyway even without the investment and the marketing.

But I want to try to (video inaudible) marketing professional can do that while appealing is simply wrong, at least in this context. When we're dealing with pharmaceuticals and disease like OUD, I think there's at least two categories of professionals who are at least as well situated, if not better, than marketing people to assess causation. And that's doctors and epidemiologists. And I want to try to explain why that is.

So I want to start with epidemiology. The Reference

Manual on Scientific Evidence tells us that epidemiology is

the field of public health and medicine that studies the incidence, distribution, and etiology of disease in human populations.

And etiology is that fancy word they always use for cause. We all want to say, "Why don't you just say cause?"

But, anyway, the point is epidemiologists, the, their, their field, their expertise of three things, incidence, distribution, cause of disease in populations. So studying causes of disease is the expertise of an epidemiologist.

Is the epidemiologist an expert in every disease? No.

Is the epidemiologist an expert in every cause independently? Again, no. But the epidemiologist's expertise is in assessing the relationship between cause and effect, of looking at the data of, you know, of an input and exposure and an effect and drilling into is that causal. That is exactly what epidemiologists do.

And I want to make a little bit of a digression. I hope I won't take up too much of your time with this. But I think it's a helpful illustration.

There was a book out a couple of years ago. I don't know if, you know, if Your Honor would have noticed it, about a guy named John Snow. And John Snow was a doctor in England and he's considered one of the founders of epidemiology.

And that's because there was a cholera pandemic in the

middle of the 19th Century. It was worldwide. It was a pandemic kind of like what we're dealing with now except it was cholera which I think is way worse.

And there was a particular outbreak in London that caught his attention. And nobody knew then what caused cholera, but Dr. Show figured out that the cholera was transmitted through contaminated water and, in particular, through a particular pump in London where a lot of people got their water.

Now, he wasn't an expert in pumps but what he was an expert in, besides medicine generally, was analyzing data in a systematic way to figure out what the cause of an outbreak was.

So he looked at who got the disease and who didn't, where it happened and where it didn't. And he made the connections to figure out, you know, based on those, those factors was this causal or not.

And since then, epidemiology has developed. It's advanced. They've come up with more advanced systems to assess whether an association is causal or not.

This is Dr. Keyes' field. She's an epidemiologist.

And not only is she an epidemiologist, her field of expertise is substance abuse. She studies the incidence, distribution, and causes of substance abuse. That's what she does when she's not working for us. That's her day job.

It has nothing to do with us having hired her.

Now, if you want to figure out how to prevent people from getting OUD, you need to understand the causes. If you don't know why we're having this outbreak, you can't figure out how to stop it. You can't prevent something if you don't know what's causing it.

And as part of her work as an epidemiologist, again not part of this case, just part of her regular job, she is part of a large NIH grant funded initiative aiming to reduce opioid overdose; specifically, you know, to develop strategies to figure out what's going to make it go away.

So here's Dr. Keyes out in the field assessing what's caused the opioid epidemic and how to prevent folks from getting it and, therefore, from overdosing.

Now, according to the defendants, Dr. Keyes may be qualified to tell us about all the other potential causes of substance abuse; familial history or mental illness or anything else that might be correlated.

But somehow, according to the defendants, when it comes to the -- this particular cause, the potential causal effect of marketing, suddenly only a marketing professional can do that. I mean, it would -- you know, it would be as if to say only a genetic counselor could assess the role of familial history in, in causing OUD.

This is what Dr. Keyes does for a living. So it's --

to say that she suddenly can address all the other potential causes using the statistical tools and sophistication, but suddenly can't talk about this one because it's marketing I don't think makes sense.

And this is particularly true because -- you know, this bleeds into the issue of methodology. The methodology here involves looking at studies. And one of the things that an epidemiologist is an expert in is reading studies of those kinds of studies, population studies, studies between cause and effect. This is what she does.

So when Dr. Keyes tells you, "Well, what I did is to read some studies," yes, that's what she does. You know, the epidemiologist looks at the studies that found correlation and figures out if it's causal.

So there is literature on the role of marketing and prescribing. And, so, there's nothing surprising about an epidemiologist looking at that.

The situation for the medical doctors, Dr. Lembke and Dr. Kolodny, is a little different, although it's similar. And, again, it comes down to the specifics of what the marketing case here is about.

So what we've said and we're going to prove are that before the defendants and the manufacturers got busy with all that marketing, doctors prescribed opioids in a certain way, in a very limited way according to certain guidelines

and precepts that kept the prescribing low.

Second, that after all the marketing, doctors prescribed differently. Suddenly they were prescribing very loosely as if every ailment needed an opioid.

And, third, and this is very important, that much of the marketing was false and fraudulent.

So, you know, if we look at what expertise a medical doctor has with respect to those elements, and then we'll talk about why that helps us with causation, a medical doctor has the expertise to tell us how doctors used to prescribe opioids and to understand what it would take for doctors to change their minds about that; if, if you are an addiction or pain or an addiction and pain doctor, to understand why did we used to do it that way and why are we now doing it differently.

In addition to that, doctors like Dr. Lembke and Dr. Kolodny have the expertise to know what about the marketing was false. And this is absolutely critical and here's why.

If everything the defendants and the manufacturers had said about opioids was true, there were a lot of ways doctors could have come to believe that mass prescribing was a good idea. They could have figured it out themselves.

They could have observed their own patients and come to that conclusion. They could have read studies that the defendants and manufacturers had nothing to do with because

there are a lot of independent sources for things that are true.

The problem is that when something is not true, it becomes easy to trace. When somebody says something false, a particular thing and someone else down the road is repeating that exact thing, it's easier to see the influence of the liar. And why is it easier to see? Because there are no independent sources for things that are false. There are many independent sources for things that are true.

So if the defendants and the manufacturers began promoting a particular set of misrepresentations and after a while lots of doctors started behaving as if those misrepresentations were true, well, where did they get that? What were the other sources available to doctors to make them believe those things? And how did they come to believe them? What was the role of continuing medical education in getting doctors to believe those things?

An addiction specialist like Dr. Lembke and Dr. Kolodny, these doctors have the expertise to know what the doctors were doing before, what it would take for them to change their mind. They have the historical knowledge of what it did take, what actually made their colleagues change their mind. And they have the expertise to trace which of the messages were false and to see how those particular false messages came to pervade the medical community.

They also have the expertise to read the literature about the effect of marketing on prescribing. And that literature, Your Honor, includes an understanding that doctors themselves are not the best reporters of the sources of their own, you know, of their own ideas about prescribing.

And I want to say a couple of things specifically about Dr. Kolodny who was not the subject of Judge Polster's order. He wasn't an expert in the CT1.

Dr. Kolodny has spent approximately 17 years researching specifically the industry-funded campaign to increase prescribing of opioids and the relationship between the opioid industry and pain organizations. He co-founded Physicians For Responsible Prescribing long before his involvement in this case which focused specifically on the role of marketing and the need for counter education to combat misinformation.

So he has spent 17 years or so devoting his life and his professional career to understanding how we got here, how did we get to this opioid crisis, and what role did the marketing of opioids play in that.

Not only that, he's been consulted by the U.S. Senate, Finance Committee specifically, on the question of the role of manufacturers and marketing in causing the opioid crisis.

He's been asked to provide his expertise about the

causes of the opioid epidemic to numerous groups, including The World Health Organization, The National Governors Association, The National Association of State Attorneys General, The National Judicial Opioid Task Force, The National Academy of Sciences, and bipartisan members of Congress. He's been invited to testify before Congress before a host of committees. And the expertise for which he's been consulted has specifically included the causes of the opioid epidemic. 

So to say that he's not qualified to talk about this one particular cause, the role of marketing in causing it, I think is really to turn on its head the notion of (video inaudible) for everyone else on the role of marketing in the opioid epidemic to say that he's not qualified to be an expert in a bench trial before Your Honor when The World Health Organization wants to hear from him on how marketing related to, to causes I think is really to stretch Daubert beyond what, what it can do.

I've taken up a lot of time with that and I want to try to be brief on my last couple of points.

The defendants say that the experts used no methodology at all. And I think that's, that's simply not correct. As we've talked about, Dr. Lembke did her own independent research. She read a lot of -- she read literature.

Dr. Kolodny has done a lot of his own independent

research on literature in delving into where did this problem come from.

Dr. Keyes used her epidemiological methods to look at the limited amount of studies that there are on this.

So I think we do have a methodology. It's just not the methodology that the defendants want, although, as I say, with Dr. Lembke, she actually did talk to doctors. But the notion that the best way to do this is to ask doctors what affected them I think is fundamentally wrong.

The case is about public health. It's about large overall risks. Not every doctor had to be convinced in order for this to become a problem. It was enough to get a core of people.

And, so, I think -- and if we did have particular interviews, then the defendants could just say, "Well, that's anecdotal. This doctor says, 'Yeah, I listened. This doctor says, 'I didn't.'"

Instead what we've looked at are the broad trends.

What were the marketing materials? Where did they do? What happened to the sales? What happened to continuing medical education? What happened to the guidelines? And was any of this true?

I want to say two things about the *Actos* decision in particular which was mentioned. I think *Actos* is quite distinguishable. I think any case where we're dealing with

individual personal injury cases requires a plaintiff to show that his or her particular doctor relied. But because we are talking about public health, we just need to show that the trend of the increase in sales that devastated the communities, that was caused by the marketing. And, so, it doesn't matter to identify particular people in order to do that.

I think for the same reason the opinions don't need to be specific to West Virginia. The marketing campaign of the defendants and the manufacturers was nationwide in scope. It aimed to change the understanding of the risks and benefits of opioids throughout the United States. And that understanding was changed, and the number of prescriptions increased substantially in West Virginia as in other parts of the country.

And these experts have studied that nationwide change and they've concluded that the marketing behavior caused the increase. And surely the fact finder could conclude that when you have the same behavior and the same outcome all over the country, and the experts have seen a causal connection overall, that it's causal here too.

The risks and benefits of opioids are the same here as everywhere, and the marketing campaign was no less fraudulent here than anywhere else.

So if somebody wants to come up with a theory that

while doctors all over the country came to believe things that weren't true because of the defendants and the manufacturers, but somehow in West Virginia doctors came to believe the same untrue things by a completely different route, I don't think it's plaintiffs' burden to eliminate that remaining speculative possibility. It may be the subject of cross-examination. It may be something for Your Honor to take into account as to weight. But I don't think it's our burden to chase down every last, you know, possibility that somehow West Virginia was on a different planet than everybody else.

I think the last thing I wanted to say -- I think I've already talked about Dr. Mohr and why the rest of her opinion is helpful to the fact finder. We understand that Dr. Mohr's testimony will need to comply with the Court's April 8th order which defendants mentioned.

But she has a lot of helpful information in understanding what the defendants were doing, how it dovetailed with the manufacturers, how manufacturers and distributors worked together, how issue management plays a role in marketing, how channel management plays a role in marketing. And I think that all of those things mean that she has many helpful opinions to provide.

To the extent that she -- that plaintiffs try to go beyond that and have her, use her to tell things that she's

not helpful for, I think the Court can curtail that at trial.

The fact that she doesn't have an opinion that any of the marketing was false is not a weakness. We're proving that through other experts. Again, it points to the limits of what a marketing expert is good at and what a marketing expert is not. We have other evidence for that.

I think, Your Honor, that unless you have further questions, I think that takes me to the end.

THE COURT: I don't. We've got a little bit of time left and I want to give Ms. Watterson the chance to respond.

Can you do it in ten minutes?

MS. WATTERSON: I can, Your Honor. Can you hear me? Did I turn on my mic? I am going to bring to bear the skills that I use in the appellate court when I only have five minutes for rebuttal. So I have a few points. It's not going to be perfect prose, but I'm going to check the box, boxes of what's on my list.

And I'm going to try to bring us back to *Daubert*, not an overall argument on the marketing case and the strength and weaknesses of the evidence. I just want to look at these experts.

Your Honor, I'm not going to say anymore during this argument about the lack of qualifications when it comes

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to -- and I don't mean to be disrespectful just using their
last names -- Keyes, Lembke and Kolodny when it comes
specifically to the issue of whether they can say marketing
drove prescribing of opioids in a way that led to
oversupply.
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I know they have opinions in other areas. That's what Judge Polster correctly said that they cannot testify on because they do not have the expertise. So that's in our brief. And again, Your Honor, we don't just rely on Judge Polster. We describe why he was correct on that.

On methodology, these experts, the three that purport to offer an opinion in this case -- again, this is plaintiffs' marketing liability here, distributors marketing, distributors marketing, whether on their own or in collaboration -- but, by the way, there is no evidence of collaboration with respect to the assertions in the complaint, the seven supposed falsehoods that drove overprescribing.

But setting that aside, there was no method brought to bear by Keyes, Kolodny, or Lembke that answers that question based upon the review of what needs to be reviewed, not just academic literature that says, well, it's possible that this could impact this.

They admitted that they did not look -- and we have citations in our brief to support, and I'm sorry I don't

have the page for you, Your Honor. They did not -- they do not know whether any distributors disseminated any marketing material. They don't know whether any doctors at all, much less those in Cabell County, saw any such marketing materials or whether any doctors relied on those materials in writing an opioid prescription.

So in order to proffer an opinion that distributors engaged in marketing that drove overprescribing, one would think you would need to know those things. Did we, in fact, disseminate material? Did anybody see such material, and whether doctors relied on those materials? That's the type of study and analysis that needs to be done, not the literature review that was discussed. And by these experts' own admissions, they didn't do that.

And I think Ms. Bierstein basically admitted because she didn't defend the notion that they looked into Cabell and Huntington. They didn't look in Cabell or Huntington. So that -- and, again, we laid this forth in our brief. They didn't engage in the right methodology.

Let me say something again about Mohr. You know, there was a suggestion that our argument on Mohr was a late-breaking argument with respect to Mohr. It was really how all of the pieces fit together, you know, given what she had disclaimed in terms of, "I don't have anything to say about falsehoods. I don't have anything to say about this

causation question." My question was limited to whether distributors engaged in marketing.

And I think that this Court's April 8th opinion, you know, speaks to that notion of, you know, Mohr getting up on the stand and just going through the litany of inventory of activities that she says are marketing. That's not helpful to the Court.

They've got to put into putting evidence saying what we did, and then the fact finder, Your Honor, needs to determine whether that, that moved the needle. And we don't need her to give a narrative on that evidence.

So, you know, just about everything that I said in this rebuttal, Your Honor, is in the brief. It is supported by citations to the record, citations to the record of these experts who were very candid in their deposition in terms of telling us what they did and didn't do, what they reviewed, and what they didn't review.

You know, Dr. Keyes, for instance -- I know I keep focusing on this because there is this assumption, not yet proven, that there was this collaboration with manufacturers. There's no evidence of that.

And Keyes herself said she didn't -- she's not really talking about distributors. She says the materials that I looked at, and these are my words, not hers, that she thinks moved the needle were manufacturer materials. And, you

know, that's in her deposition. We've cited to those pages in our brief.

So, again, taking it back to Daubert, I'm not going to respond on all of the arguments with respect to the marketing theory, but we have three experts who may be highly qualified to testify in other areas and may very well be on the stand in this case. They simply are not qualified to testify with respect to the marketing theory.

Jakki Mohr has nothing to say on causation or falsity.

And all she's really going to provide to the Court is a laundry list and inventory.

And I'm arguing on qualifications, not asking you to rule the same way Judge Polster ruled just because he ruled that way but, instead, because that is the correct outcome here.

One last comment on that. There was an assertion -and, again, I'm doing a checklist here, not in a beautiful
narrative. There was a suggestion that something had
changed between the time that Judge Polster issued his
opinion and today, a suggestion that some of the arguments
and reasons why plaintiffs believe Lembke and Keyes are
qualified were either not before Judge Polster or not
understood by Judge Polster.

We explained in our brief, again with support, why that is incorrect. Judge Polster did have in front of him

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     everything that Your Honor has in front of you today.
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          And thank you for taking so much time on this marketing
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     issue, or giving the parties so much time. Appreciate it.
               THE COURT: Well, I realize that it's important
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     for counsel to know what the result here is and I will --
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     all I can promise you is that I'll rule on these things as
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     promptly as I can. And I thank you all very much. These
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     are very helpful arguments.
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          (Proceedings concluded at 11:58 a.m.)
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1	I, Lisa A. Cook, Official Reporter of the United
2	States District Court for the Southern District of West
3	Virginia, do hereby certify that the foregoing is a true and
4	correct transcript, to the best of my ability, from the
5	record of proceedings in the above-entitled matter.
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7	
8	s\Lisa A. Cook <u>April 15, 2021</u>
9	Reporter Date
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